MEMORANDUM

TO: Mr. Addison Rice

Anderson, Mulholland and Associates

DATE: June 27, 2016

FROM: R. Infante

FILE: 1606028C

RE:

Data Validation Air samples SDG: 1606028C

SUMMARY

Full validation was performed on the data for several gas samples analyzed for methanol by Compendium Method TO-15. Determination of Volatile Organic Compounds (VOCs) In Air Collected In Specially-Prepared Canisters and Analyzed By Gas Chromatography/Mass Spectrometry (GC/MS), January, 1999". The samples were collected at the Bristol Myer Squib facility, Humacao, PR site on May 29, 2016 and submitted to Eurofins Air Toxics, Inc. of Folson, California that analyzed and reported the results under delivery groups (SDG) 1606028C.

The sample results were assessed according to USEPA data validation guidance documents in the following order of precedence: Volatile Organic Analysis of Ambient Air in Canisters by Method TO-15, (SOP # HW-31. Revision #4. October, 2006. The QC criteria and data validation actions listed on the data review worksheets are from the primary guidance document, unless otherwise noted.

In general the data is valid as reported and may be used for decision making purposes. The data results are acceptable for use.

SAMPLES

The samples included in the review are listed below

Client Sample ID	Lab. Sample ID	Collected Date	Matrix	Analysis
B30AA (052816)	1606028C-01A	05/29/2016	Air	Methanol
B30IA-1 (052816)	1606028C-02A	05/29/2016	Air	Methanol
B30IA-1D (052816)	1606028C-03A	05/29/2016	Air	Methanol
B30IA-2 (052816)	1606028C-04A	05/29/2016	Air	Methanol
B30IA-3 (052816)	1606028C-05A	05/29/2016	Air	Methanol
B30IA-4 (052816)	1606028C-06A	05/29/2016	Air	Methanol
B30IA-5 (052816)	1606028C-07A	05/29/2016	Air	Methanol

REVIEW ELEMENTS

Sample data were reviewed for the following parameters, where applicable to the method

- o Agreement of analysis conducted with chain of custody (COC) form
- o Holding time and sample preservation
- Gas chromatography/mass spectrometry (GC/MS) tunes
- o Initial and continuing calibrations
- o Method blanks/trip blanks/field blank
- o Canister cleaning certification criteria
- Surrogate spike recovery
- o Internal standard performance and retention times
- o Field duplicate results
- o Laboratory control sample/laboratory control sample duplicate (LCS/LCSD) results
- o Quantitation limits and sample results

DISCUSSION

Agreement of Analysis Conducted with COC Request

Sample reports corresponded to the analytical request designated on the chain-of-custody form.

Holding Times and Sample Preservation

All samples analyzed within the recommended method holding time. All summa canisters received in good conditions except for the cases described in this document. The Summa canister for sample B30IA-2 (052816) was leaking upon arrival. The client was notified and the analysis proceeded. Reported analyte concentrations were qualified by the laboratory as estimated. No further qualification made.

Samples analyzed within method recommended holding time.

Initial and Continuing Calibrations

Methanol by Compendium Method TO-15

One point calibration performed. Initial and continuing calibrations meet method specific requirements. Initial calibration retention times meet method specific requirements.

Method Blank/Trip Blank/Field Blank

Target analytes were not detected in laboratory method blanks.

No trip/field blank analyzed with this data package.

Laboratory/Field Duplicate Results

Field/laboratory duplicates were analyzed as part of this data set. Target analytes meet the RPD performance criteria of +25% for analytes $5\times SQL$.

LCS/LCSD Results

Methanol

No LCS/LCSD (blank spike) was analyzed by the laboratory associated with this data package.

Quantitation Limits and Sample Results

Dilutions were performed (see worksheet).

Calculations were spot checked.

Certification

The following samples 1606028C-01A; 1606028C-02A; 1606028C-03A; 1606028C-04A; 1606028C-05A; 1606028C-06A; and 1606028C-07A were analyzed following standard procedures accepted by regulatory agencies. The quality control requirements met the methods criteria except in the occasions described in this document.

Rafael Infante

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Client Sample ID: B30AA (052816) Lab ID#: 1606028C-01A

File Name: Dil. Factor:	14061304 2.30		Date of Collection: 5/29/16 6:24:00 PM Date of Analysis: 6/13/16 10:17 AM			
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)		
Methanol	120	Not Detected	150	Not Detected		
Container Type: 6 Liter Summ	na Canister (100% Certifie	d)				
Surrogates		%Recovery		Method Limits		
1,2-Dichloroethane-d4		107		70-130		
Toluene-d8		100		70-130		
4-Bromofluorobenzene		102		70-130		





Client Sample ID: B30IA-1 (052816) Lab ID#: 1606028C-02A

File Name: Dil. Factor:	14061305 1.56		Date of Collection: 5/29/16 5:39:00 PN Date of Analysis: 6/13/16 10:35 AM			
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt Limit (ug/m3)	Amount (ug/m3)		
Methanol	78	Not Detected	100	Not Detected		
		44				
Container Type: 6 Liter Sumn	na Canister (100% Certifie	d)				
Container Type: 6 Liter Sumn Surrogates	na Canister (100% Certifie	%Recovery		Method Limits		
	na Canister (100% Certifie	•				
Surrogates	na Canister (100% Certifie	%Recovery		Limits		





Client Sample ID: B30IA-1D (052816) Lab ID#: 1606028C-03A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	14061306 1.56	Date of Collection: 5/29/16 5:39:00 PM Date of Analysis: 6/13/16 10:54 AM			
	Rpt. Limit	Amount	Rpt. Limit	Amount	
Compound	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)	
Methanol	78	Not Detected	100	Not Detected	

Container Type: 6 Liter Summa Canister (100% Certified)

Surrogates	%Recovery	Method Limits
1,2-Dichloroethane-d4	104	70-130
Toluene-d8	100	70-130
4-Bromofluorobenzene	105	70-130





Client Sample ID: B301A-2 (052816) Lab ID#: 1606028C-04A

File Name: Dil. Factor:	14061307 1.32	Date of Collection: 5/29/16 10:45:00 AM Date of Analysis: 6/13/16 11:23 AM			
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)	
Methanol	66	Not Detected	86	Not Detected	
Container Type: 6 Liter Summ	na Canister (100% Certifie	d)			
Surrogates_		%Recovery		Method Limits	
		%Recovery			
Surrogates 1,2-Dichloroethane-d4 Toluene-d8	<u>, , , , , , , , , , , , , , , , , , , </u>			Limits	





Client Sample ID: B30IA-3 (052816) Lab ID#: 1606028C-05A

File Name: Dil. Factor:	14061308 1.52	Date of Collection: 5/29/16 6:00:00 PM Date of Analysis: 6/13/16 11:42 AM			
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt Limit (ug/m3)	Amount (ug/m3)	
Methanol	76	Not Detected	100	Not Detected	
Container Type: 6 Liter Sumr	na Canister (100% Certifie	d)			
	na Canister (100% Certifie	d) %Recovery		Method Limits	
Surrogates	na Canister (100% Certifie	•			
Container Type: 6 Liter Sumr Surrogates 1,2-Dichloroethane-d4 Toluene-d8	na Canister (100% Certifie	%Recovery	Λ.	Limits	





Client Sample ID: B30IA-4 (052816) Lab ID#: 1606028C-06A

File Name: Dil. Factor:	14061309 1.76	Date of Collection: 5/29/16 6:04:00 PM Date of Analysis: 6/13/16 12:42 PM			
Compound	Rpt. Limit (ppbv)	Amount Rpt. Limit (ppbv) (ug/m3)		Amount (ug/m3)	
Methanol	88	Not Detected	120	Not Detected	
Container Type: 6 Liter Summ	na Canister (100% Certifie	d)			
				Method	
Surrogates		%Recovery		Limits	
1,2-Dichloroethane-d4		106		70-130	
Toluene-d8		99		70-130	
4-Bromofluorobenzene		104		70-130	





Client Sample ID: B30IA-5 (052816) Lab ID#: 1606028C-07A

File Name: Dil. Factor:	14061310 1.53		Date of Collection: 5/29/16 5:36:00 PM Date of Analysis: 6/13/16 01:07 PM			
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)		
Methanol	76	Not Detected	100	Not Detected		
Container Type: 6 Liter Sumn	na Canister (100% Certifie	d)				
	na Canister (100% Certifie	d) %Recovery		Method Limits		
Container Type: 6 Liter Sumn Surrogates 1,2-Dichloroethane-d4	na Canister (100% Certifie	•				
Surrogates	na Canister (100% Certifie	%Recovery		Limits		



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Sample Transportation Notice
Relinquishing signature on this document indicates that sample is being shipped in compliance with all applicable local, State, Federal, national, and international laws, regulations and ordinances of any kind. Air Toxics Limited assumes no liability with respect to the collection, handling or shipping of these samples. Refinquishing signature also indicates agreement to hold harmless, defend, and indemnify Air Toxics Limited against any claim, demand, or action, of any kind, related to the

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Lab 1.0. Field Sample I.D. (Location)	Can #	_	laction	Time of Collection	Analyses Design	-44			ssure/Vac	
A very August 5 feb	2 2 7 1 1 1	-		OF CORECTION	Analyses Reque	sted	Initial	Final	Receipt	Final (psi)
B30 AA (052816)	33321	5/2	9/16	1824	70-15		301	7.5		
1031 B30IA-1(052816)	35258	5/2	29/16	1739	To-15	·	30°	5.5		
55A B30IA-10(052816)	14038	5/1	19/16	1739	TO-15		29	5,0		
044 B30 A-2 (052816)	33586				TO-15		30	0		
054 B30IA-3(052816)	33926	7.	7.	1800	TO-15			4		100000
00 B30TA-4 (052816)	0337						29			
07A B30IA-5 (052816)					TO-15		30	8	13 12 A	
0301A 5 (031g(C)	6L1279	5/2	29/16	1736	TO-15		28.5	4		
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	Project Number:1606028C Date:05/29/2016
REVIEW OF VOLATILE ORGANIC Parameters for evaluating volatile organics were creations. This document will assist the reviewer in using profession decision and in better serving the needs of the data users. The samp USEPA data validation guidance documents in the following ord "Compendium Method TO-15. Determination of Volatile Organic Compendium Method TO-15. Determination of Volatile Organic Compendium Method TO-15. Determination of Volatile Organic Compendium, 1999"; USEPA Hazardous Waste Support Branch. Validanalysis of Ambient Air in Canisters by Method TO-15, (SOP # HW-QC criteria and data validation actions listed on the data review work document, unless otherwise noted. The hardcopied (laboratory name) _EurofinsAir_Toxics	ated to delineate required validation all judgment to make more informed all results were assessed according to the results were assess
Lab. Project/SDG No.:1606028C No. of Samples:7	Sample matrix:Air
Trip blank No.: Field blank No.: Equipment blank No.: Field duplicate No.:B30IA-1_(052816)/_B30IA-1D_(05281	
X Holding TimesXX GC/MS TuningXX Internal Standard PerformanceXX BlanksXX Surrogate RecoveriesXN/A_ Matrix Spike/Matrix Spike Duplicate	Laboratory Control Spikes Field Duplicates Calibrations Compound Identifications Compound Quantitation Quantitation Limits
Overall Comments:Methanol_by_method_TO-15	
Definition of Qualifiers: J- Estimated results U- Compound not detected R- Rejected data UJ- Estimated nondetect Reviewer: Lafaul Maux	<u> </u>
Date:06/27/2016	

DATA REVIEW WORKSHEETS

DATA COMPLETENESS

MISSING INFORMATION	DATE LAB. CONTACTED	DATE RECEIVED
	3	
1		
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		<u></u>
1		<u> </u>
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	**	

All criteria were met _	х_
Criteria were not met	
and/or see below	

HOLDING TIMES

The objective of this parameter is to ascertain the validity of the results based on the holding time of the sample from time of collection to the time of analysis.

Complete table for all samples and note the analysis and/or preservation not within criteria

SAMPLE ID	DATE SAMPLED	DATE ANALYZED	рH	ACTION
		,		
				summa canisters received
				. The Summa canister for
sample B30IA-2	(052816) was leaking	ng upon arrival. The cl	ient was	notified and the analysis
proceeded. Repo	orted analyte concent	rations were qualified b	y the la	boratory as estimated. No
further qualification		•		•
		,		
	,			
	T			

Criteria

Aqueous samples – 14 days from sample collection for preserved samples (pH \leq 2, 4°C), no air bubbles.

Aqueous samples – 7 days from sample collection for unpreserved samples, 4°C, no air bubbles. Soil samples- 7 days from sample collection.

Cooler temperature (Criteria: 4 ± 2 °C): N/A – summa canisters

Actions

If the VOCs vial(s) have air bubbles, estimate positive results (J) and reject nondetects (R).

If the % solids of soil samples is 10-50%, estimates positive results (J) and nondetects (UJ)

If the % solid of soil samples is < 10%, estimate positive results (J) and reject nondetects (R).

If holding times are exceeded but < 14 days beyond criteria, estimate positive results (J) and nondetects (UJ).

If holding times are exceeded but < 28 days beyond criteria, estimate positive results (J) and reject nondetects (R).

If holding times are grossly exceeded (> 28 days beyond criteria), reject all results (R).

If samples were not iced or if the ice were melted (> 10°C), estimate positive results (J) and nondetects (UJ).

All criteria were met _X
Criteria were not met see below

GC/MS TUNING

The assessment standard tuning (_	determine if the sample instrume	ntation is within the
_XThe BFE	performance results were r	reviewed and found to be within the	e specified criteria.
XBFB tuni	ing was performed for every	24 hours of sample analysis.	
If no, use profes qualified or rejec		ne whether the associated data s	should be accepted,
List	the	samples	affected:

If mass calibration is in error, all associated data are rejected.

All criteria were met _	Х
Criteria were not met	
and/or see below	

CALIBRATION VERIFICATION

Compliance requirements for satisfactory instrument calibration are established to ensure that the instrument is capable of producing and maintaining acceptable quantitative data.

Date of initial calibration:	06/13/2016
Dates of continuing calibration	:06/13/2016
Instrument ID numbers:M	ISD-14
Matrix/Level:	Air/low

DATE	LAB ID#	FILE	CRITERIA OUT RFs, %RSD, %D, r	COMPOUND	SAMPLES AFFECTED
	1				
			and continuing calibrates the continuing calibrates and c	ions meet method specificulirements.	requirements. Initial
	-				

Criteria

All RFs must be > 0.05 regardless of method requirements for SPCC.

All %RSD must be \leq 15 % regardless of method requirements for CCC.

All %Ds must be \leq 30% regardless of method requirements for CCC.

Method TO-15 does not specify criterion for the curve correlation coefficient (r). A limit for r of \geq 0.995 has therefore been utilized as professional judgment.

Actions

If any compound has an initial RF or a continuing RF of < 0.05, estimate positive results (J) and reject nondetects (R), regardless of method requirements.

If any compound has a %RSD > 15%, estimate positive results (J) and use professional judgment to qualify nondetects.

If any compound has a %RSD > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and nondetects (UJ).

If any compound has a % D > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has r < 0.995, estimate positive results and nondetects.

A separate worksheet should be filled for each initial curve

All criteria were met _	Х
Criteria were not met	
and/or see below	

V A. BLANK ANALYSIS RESULTS (Sections 1 & 2)

The assessment of the blank analysis results is to determine the existence and magnitude of contamination problems. The criteria for evaluation of blanks apply only to blanks associated with the samples, including trip, equipment, and laboratory blanks. If problems with any blanks exist, all data associated with the case must be carefully evaluated to determine whether or not there is an inherent variability in the data for the case, or if the problem is an isolated occurrence not affecting other data.

List the contamination in the blanks below. High and low levels blanks must be treated separately.

Laboratory blanks

DATE ANALYZED	LAB ID	LEVEL/ MATRIX	COMPOUND	CONCENTRATION UNITS
All_method	2000 890	87 SON 7		
Field/Equipment		LEVEL/	COMPOUND	CONCENTRATION
ANALYZED No_field/trip/equ	uipment_blanks	MATRIX _analyzed_with	n_this_data_package	UNITS

All criteria were metX	
Criteria were not met	
and/or see below	

VB. BLANK ANALYSIS RESULTS (Section 3)

Blank Actions

Action Levels (ALs) should be based upon the highest concentration of contaminant determined in any blank. Do not qualify any blank with another blank. The ALs for samples which have been diluted should be corrected for the sample dilution factor and/or % moisture, where applicable. No positive sample results should be reported unless the concentration of the compound in the samples exceeds the ALs:

ALs = 10x the amount of common contaminants (methylene chloride, acetone, 2-butanone, and toluene)

ALs = 5x for any other compounds

Specific actions are as follows:

If the concentration is < sample quantitation limit (SQL) and \le AL, report the compound as not detected (U) at the SQL.

If the concentration is \geq SQL but \leq AL, report the compound as not detected (U) at the reported concentration.

If the concentration is \geq SQL and > AL, report the concentration unqualified.

Notes:

High and low level blanks must be treated separately

Compounds qualified "U" for blank contamination are still considered "hits" when qualifying for calibration criteria.

CONTAMINATION SOURCE/LEVEL	COMPOUND	CONC/UNITS	AL/UNITS	SQL	AFFECTED SAMPLES
					100
					ALCO AND
				100	

All criteria were met _	X_
Criteria were not met	
and/or see below	

SURROGATE SPIKE RECOVERIES

Laboratory performance of individual samples is established by evaluation of surrogate spike recoveries. All samples are spiked with surrogate compounds prior to sample analysis. The accuracy of the analysis is measured by the surrogate percent recovery. Since the effects of the sample matrix are frequently outside the control of the laboratory and may present relatively unique problems, the validation of data is frequently subjective and demands analytical experience and professional judgment.

List the percent recoveries (%Rs) which do not meet the criteria for surrogate recovery. Matrix: solid/aqueous

SAMPLE ID

SURROGATE COMPOUND

ACTION

1,2-DICHLOROETHANE**d4**

Toluene-

4-BFB

d8

_Surrogate_recoveries_within_laboratory_control_limits					
		-			
QC Limits* (Air)					
LL_to_UL70to_130	_70to_13070to_130				

- QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- If QC limits are not available, use limits of 80 120 % for aqueous and 70 130 % for solid samples.

Actions:

QUALITY	%R < 10%	%R = 10% - LL	%R > UL
Positive results	J	J	J
Nondetects results	R	UJ	Accept

Surrogate action should be applied:

If one or more surrogate in the VOC fraction is out of specification, but has a recovery of > 10%.

If any one surrogate in a fraction shows < 10 % recovery.

All criteria were met
Criteria were not met
and/or see belowN/A

VII. A MATRIX SPIKE/MATRIX SPIKE DUPLICATE (MS/MSD)

This data is generated to determine long term precision and accuracy in the analytical method for various matrices. This data alone cannot be used to evaluate the precision and accuracy of individual samples. If any % R in the MS or MSD falls outside the designated range, the reviewer should determine if there are matrix effects, i.e. LCS data are within the QC limits but MS/MSD data are outside QC limit.

1. MS/MSD Recoveries and Precision Criteria

The laboratory should use one MS and a duplicate analysis of an unspiked field sample if target analytes are expected in the sample. If target analytes are not expected, MS/MSD should be analyzed.

MPOUND	% R	RPD	QC LIMITS	ACTION
 t_required_as_p	art_of_N	vlethod_	TO-15;_blank_spi	ke_used_to_assess_
)	MPOUND ot_required_as_p	t_required_as_part_of_l	t_required_as_part_of_Method_	ot_required_as_part_of_Method_TO-15;_blank_spi

Actions:

QUALITY	%R < LL	%R > UL
Positive results	J	J
Nondetects results	R	Accept

MS/MSD criteria apply only to the unspiked sample, its dilutions, and the associated MS/MSD samples:

If the % R for the affected compounds were < LL (or 70 %), qualify positive results (J) and nondetects (UJ).

If the % R for the affected compounds were > UL (or 130 %), only qualify positive results (J).

If 25 % or more of all MS/MSD %R were < LL (or 70 %) or if two or more MS/MSD %Rs were < 10%, qualify all positive results (J) and reject nondetects (R).

A separate worksheet should be used for each MS/MSD pair.

^{*} QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.

^{*} If QC limits are not available, use limits of 70 – 130 %.

All criteria w	vere met	
Criteria wer	e not met	
and/or see I	below N/A	

VII. B MATRIX SPIKE/MATRIX SPIKE DUPLICATE

MS/MSD - Unspiked Compounds

It should be noted that Method TO-15 does not specify a MS/MSD criteria for the unspiked compounds in the sample. A %RSD of < 50% has therefore been utilized as professional judgment.

If all target analytes were spiked in the MS/MSD, this review element is not applicable.

List the %RSD of the compounds which do not meet the criteria.

Sample ID:			Matrix/Level/Unit			
COMPOUND	SAMPLE CONC.	MS CONC.	MSD CONC.	% RSD	ACTION	
50 S			- Con (c)			
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	-		The state of the s			
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	-					
Town						

Actions:

^{*} If the % RSD > 50, qualify the positive result in the unspiked samples as estimated (J).

^{*} If the % RSD is not calculated (NC) due to nondetected value, use professional judgment to qualify the data.

All criteria were met _X				
Criteria were not met				
and/or see below				

VIII. LABORATORY CONTROL SAMPLE (LCS) ANALYSIS

This data is generated to determine accuracy of the analytical method for various matrices.

LCS Recoveries Criteria

Where LCS spiked with the same analyte at the same concentrations as the MS/MSD? Yes or No. If no make note in data review memo.

List the %R of compounds which do not meet the criteria

COMPOUND	% R	QC LIMIT
nike) analyzed in this da	ta nackane	
	12	
		pike)_analyzed_in_this_data_package

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit
- * If QC limits are not available, use limits of 70 130 %.

Actions:

QUALITY	%R < LL	%R > UL
Positive results	J	J
Nondetects results	R	Accept

All analytes in the associated sample results are qualified for the following criteria.

If 25 % of the LCS recoveries were < LL (or 70 %), qualify all positive results (j) and reject nondetects (R).

If two or more LCS were below 10 %, qualify all positive results as (J) and reject nondetects (R).

2. Frequency Criteria:

Where LCS analyzed at the required frequency and for each matrix? Yes or <u>No</u>. If no, the data may be affected. Use professional judgment to determine the severity of the effect and qualify data accordingly. Discuss any actions below and list the samples affected.

		All criteria were metX Criteria were not met and/or see below
IX.	LABORATORY DUPLICATE PRECISION	
	Sample IDs:_B30IA-1_(052816)/_B30IA-1D_(052816)_	Matrix:Air

Field duplicates samples may be taken and analyzed as an indication of overall precision. These analyses measure both field and lab precision; therefore, the results may have more variability than laboratory duplicates which only laboratory performance. It is also expected that soil duplicate results will have a greater variance than water matrices due to difficulties associated with collecting identical field duplicate samples.

The project QAPP should be reviewed for project-specific information. Suggested criteria: RPD \pm 25% for air samples. If both samples and duplicate are <5 SQL, the RPD criteria is doubled.

COMPOUND	SQL	SAMPLE CONC.	DUPLICATE CONC.	RPD	ACTION				
RPD within the method performance criteria.									

Actions:

Qualify as estimated positive results (J) and nondetects (UJ) for the compound that exceeded the above criteria. For organics, only the sample and duplicate will be qualified.

If an RPD cannot be calculated because one or both of the sample results is not detected, the following actions apply:

If one sample result is not detected and the other is greater than 5x the SQL qualify (J/UJ).

If one sample value is not detected and the other is greater than 5x the SQL and the SQLs for the sample and duplicate are significantly different, use professional judgment to determine if qualification is appropriate.

If one sample value is not detected and the other is less than 5x, use professional judgment to determine if qualification is appropriate.

If both sample and duplicate results are not detected, no action is needed.

Actions:

All criteria were met	X
Criteria were not met	
and/or see below	

X. INTERNAL STANDARD PERFORMANCE

The assessment of the internal standard (IS) parameter is used to assist the data reviewer in determining the condition of the analytical instrumentation.

List the internal standard area of samples which do not meet the criteria.

- * Area of +40% or -40% of the IS area in the associated calibration standard.
- * Retention time (RT) within \pm 0.06 seconds of the IS area in the associated calibration standard.

DATE	SAMPLE ID	IS OUT	IS AREA	ACCEPTABLE RANGE	ACTION
	tandard_area_and_reration_standards				_both_samples
				- 110 -	
	N-331				
	2511, 04				

1. IS actions should be applied to the compound quantitated with the out-of-control ISs

QUALITY	IS AREA < -40%	IS AREA > + 40%
Positive results	J	J
Nondetected results	R	ACCEPT

 If a IS retention time varies more than 0.330 seconds, the chromatographic profile for that sample must be examined to determine if any false positive or negative exists. For shifts of a large magnitude, the reviewer may consider partial or total rejection of the data for the sample fraction.

All criteria were met __X__ Criteria were not met and/or see below ____

XII. SAMPLE QUANTITATION

The sample quantitation evaluation is to verify laboratory quantitation results. In the space below, please show a minimum of one sample calculation:

Calibration check

1,2-Dichloroethane-d4

RF = 1.76119

[] = (173195)(400)/(92008)(1.76119)

= 427.5 ppbv OK

All criteria were met _X	
Criteria were not met	
and/or see below	

XII. QUANTITATION LIMITS

A. Dilution performed

SAMPLE ID	DILUTION FACTOR	REASONS FOR DILUTION
All samples dil	uted by a factor of less th	an 2.3.
		THE REAL PROPERTY.
a)		
	- COS	

3.	Percent Solids		
	List samples which have ≤ 50 % solids		
		1000	101
			= =======

Actions:

If the % solids of a soil sample is 10-50%, estimate positive results (J) and nondetects (UJ)

If the % solids of a soil sample is < 10%, estimate positive results (J) and reject nondetects (R)